FDA 510(k) Premarket Notification

K032066 SONOSURG SYSTEM

CLEARED

Oct 3, 2003
92 days to decision

K032066 · Product code: **LFL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k032066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jul 3, 2003
Decision date	Oct 3, 2003
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Olympus Optical Co.**
Location	Melville, NY, US
Contact	TINA STEFFANIE-OAK
510(k) history	22 submissions · 22 cleared · 1998-2003